

SENTARA COMMUNITY PLAN (MEDICAID)

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-800-750-9692. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If the information provided is not complete, correct, or legible, the authorization process can be delayed.

Drug Requested: Spravato[®] (esketamine)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight: _____ Date: _____

Quantity Limit:

- Major Depressive Disorder with Acute Suicidal Ideation or Behavior: 8 kits/month; 1 month of treatment
- Treatment-Resistant Depression: 4 kits/month (*induction dose requires 8 kits/month)

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Choose **ONE** of the following applicable diagnoses below. **Provider Please Note:** Any indication that is **NOT** FDA approved will be considered experimental/investigational and **NOT** medically necessary

Treatment Resistant Depression. ALL the following criteria must be met:

Length of Authorization: 1 year

(Continued on next page)

- Member must be 18 years of age or older

AND

- Spravato[®] must be prescribed by or in consultation with a psychiatrist
 - Provider is a psychiatrist
 - Consult with psychiatrist (include name/date): _____

AND

- Member must have a diagnosis of treatment resistant depression (TRD) without psychotic features defined by current DSM criteria made or verified by a psychiatrist
 - ICD Code/Diagnosis: _____

AND

- Member must be experiencing moderate to severe symptomology documented by a standardized rating scale that reliably measures depressive symptoms. **A current baseline (within previous 30 days, prior to starting Spravato[®]) scale with scoring must be attached.**
 - Scale: _____
 - Date Administered: _____

AND

- Member must have experienced clinical failure or intolerance with at least two (2) antidepressant therapies from at least two (2) different drug classes (**verified by pharmacy paid claims and/or chart notes**)
 - Failures must be of adequate dose (maximally tolerated)
 - Failures must be of adequate duration (at least 6 weeks)
 - Adherent fills required (verified by pharmacy claims)
 - Failures must occur during current depressive episode
 - Antidepressant therapy would include any of the following classes:
 - Selective serotonin reuptake inhibitors (e.g., citalopram, fluoxetine, paroxetine, sertraline)
 - Serotonin norepinephrine reuptake inhibitors (e.g., duloxetine, venlafaxine)
 - Bupropion
 - Tricyclic antidepressants (e.g., amitriptyline, clomipramine, nortriptyline)
 - Mirtazapine
 - Monoamine oxidase inhibitors (e.g., selegiline, tranylcypromine)
 - Serotonin modulators (e.g., nefazodone, trazodone)

1. Drug: _____ Dose: _____ Duration: _____

Reason for Discontinuation: _____

2. Drug: _____ Dose: _____ Duration: _____

Reason for Discontinuation: _____

AND

- Member must have experienced clinical failure or intolerance with at least one (1) augmentation therapy (e.g., lithium, liothyronine, antipsychotics or anticonvulsants) (**verified by pharmacy paid claims and/or chart notes**)

- Failures must be of adequate dose (maximally tolerated)
- Failures must be of adequate duration (at least 6 weeks)
- Adherent fills required (verified by pharmacy claims)
- Failures must occur during current depressive episode

1. Drug: _____ Dose: _____ Duration: _____

Reason for Discontinuation: _____

2. Drug: _____ Dose: _____ Duration: _____

Reason for Discontinuation: _____

AND

- Spravato[®] must be used in combination with a newly initiated daily oral antidepressant that has not been previously tried. **Documentation (pharmacy claims or chart notes) required.**

Drug: _____

AND

- Member does **NOT** have aneurysmal vascular disease (including thoracic and abdominal aorta, intracranial and peripheral arterial vessels), arteriovenous malformation, or a history of intracerebral hemorrhage

AND

- Prescriber must have assessed the member's risk for abuse of controlled substances (i.e., review of medical history, review of state prescription monitoring program (PMP))

AND

- Member must be enrolled in the Spravato[®] REMS program

AND

- Administering site/provider must be certified in the Spravato[®] REMS program:

Name/Location of Administering Provider: _____

Diagnosis: Major Depressive Disorder with Suicidal Ideation or Behavior

Continuation of inpatient Spravato[®] therapy, ALL the following criteria must be met:

**One-time authorization per episode for remaining doses required for continuation.
Maximum allowable duration = 1 month**

- Provider **MUST** submit date of therapy initiation and number of doses administered up to point of request

Date Spravato[®] therapy initiated: _____

Number of doses administered since initiation: _____

AND

- Member must be 18 years of age or older

Diagnosis: Major Depressive Disorder with Suicidal Ideation or Behavior

Initiation of outpatient Spravato® therapy, ALL the following criteria must be met:

One-time authorization per episode for a duration of 1 month, total of 8 kits/month

- Member must be 18 years of age or older

AND

- Spravato® must be prescribed by or in consultation with a psychiatrist

- Provider is a psychiatrist

- Consult with psychiatrist (include name/date): _____

AND

- Member must have a diagnosis of major depressive disorder with acute suicidal ideation or behavior verified by a psychiatrist.

AND

- Spravato® must be used in combination with a daily oral antidepressant. Documentation (pharmacy claims or chart notes) required.

- Drug: _____

AND

- Prescriber must have assessed the member's risk for abuse of controlled substances (i.e., review of medical history, review of state prescription monitoring program (PMP))

AND

- Member must be enrolled in the Spravato® REMS program

AND

- Administering site/provider must be certified in the Spravato® REMS program:

- Name/Location of Administering Provider: _____

Medication being provided by Specialty Pharmacy – Proprium Rx

*****Use of samples to initiate therapy does not meet step edit/preauthorization criteria.*****
****Previous therapies will be verified through pharmacy paid claims or submitted chart notes.****